RWANDA FDA
Rwanda Food and Drugs Authority

Kigali on 14/03/2024
Ref. N°: FDISM/FDIC/1267./FDA/2024

All Stakeholders

RE: Circular on public consultation and call for comments on regulatory documents

Reference is made to the Law N° 003/2018 of 09/02/2018 establishing Rwanda Food and Drugs Authority (Rwanda FDA) and determining its mission, organization and functioning especially in its article 8 paragraph 2, the Authority regulates compliance with quality standards relating to the manufacture, storage, sale, distribution, use, import and export, labels, packages and raw materials used in the manufacture of products regulated under this Law.

Rwanda FDA would like to inform all stakeholders and the general public that the draft of updated Regulations governing licensing of public and private manufacturers, distributors, wholesalers and retailers of medical Products, Doc. No. FDISM/FDIC/TRG/001; Regulations Governing Good Storage and Good Distribution Practices of Medical products, Doc. No. FDISM/FDIC/TRG/006; Guidelines for Good Storage and Distribution Practices of Medical Products, Doc. No. FDSIM/FDIC/GDL/006, is being circulated for public consultation and comments starting from 12th March 2024

These guidelines have been uploaded to Rwanda FDA website https://rwandafda.gov.rw/ under stakeholders rubric and will be available for 30 calendar days. The Authority invites all stakeholders to provide inputs/comments using comment form No ODG/QMS/FMT/023 and send the completed form to the email: info@rwandafda.gov.rw not letter than 11/04/2024.Thereafter, there will be a stakeholder meeting to validate the inputs/comments.

Sincerely,

Prof. Emile BIENVENU
Director General

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